## DAX1 (D2F1) Rabbit mAb



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## For Research Use Only. Not for Use in Diagnostic Procedures.

<b>Applications:</b> WB, IP, IHC-P, IF-IC	Reactivity: H	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 48	Source/Isotype: Rabbit IgG	UniProt ID: #P51843	Entrez-Gene Id 190
Product Usage Information	Ар	plication				Dilution
	We	estern Blotting				1:1000
	lmi	munoprecipitation				1:100
	lmi	Immunohistochemistry (Paraffin)				1:400
	lmi	Immunofluorescence (Immunocytochemistry)				1:3200
Storage	•	Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 $\mu$ g/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.				
Specificity / Sensit	ivity DAX	DAX1 (D2F1) recognizes endogenous levels of total DAX1 protein.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Gly330 of human DAX1 protein.				
Background	the binc tran pitui prod mal mai mut	DSS-AHC critical region on the X chromosome protein 1 (DAX1) is an orphan nuclear receptor encoded by the nuclear receptor subfamily 0 group B member 1 ( <i>NR0B1</i> ) gene. DAX1 possesses an atypical DNA binding domain that allows it to form heterodimeric complexes with DNA binding partners and repress transcriptional activity (1,2). During development, DAX1 is important for establishment of the hypothalamic-pituitary-adrenal gonadal axis. The receptor is essential for development of several important hormone-producing organs that determine this axis, including the adrenal glands, pituitary, hypothalamus, and the male and female reproductive organs (3,4). Research studies suggest that DAX1 plays a role in maintenance of pluripotency in embryonic stem cells (5,6). Loss of DAX1 function through deletion or mutation results in adrenal insufficiency and hypogonadotropic hypogonadism (7), while duplication of the <i>NR0B1</i> gene on the X-chromosome causes dosage-sensitive sex reversal (8).				
Background Refere	2. ly 3. N 4. M 5. U 6. W 7. J	<ol> <li>Iyer, A.K. et al. (2006) Mol Endocrinol 20, 2326-42.</li> <li>Iyer, A.K. and McCabe, E.R. (2004) Mol Genet Metab 83, 60-73.</li> <li>Niakan, K.K. and McCabe, E.R. (2005) Mol Genet Metab 86, 70-83.</li> <li>McCabe, E.R. (2007) Mol Cell Endocrinol 265-266, 179-82.</li> <li>Uranishi, K. et al. (2013) Mol Cell Biol 33, 2056-66.</li> <li>Wang, Q. and Cooney, A.J. (2013) Adv Exp Med Biol 786, 287-306.</li> <li>Jadhav, U. et al. (2011) Mol Cell Endocrinol 346, 65-73.</li> <li>Sanlaville, D. et al. (2004) Am J Med Genet A 128A, 325-30.</li> </ol>				

Species reactivity is determined by testing in at least one approved application (e.g., western blot). **Species Reactivity** 

Western Blot Buffer IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X TBS,

0.1% Tween® 20 at 4°C with gentle shaking, overnight.

WB: Western Blotting IP: Immunoprecipitation IHC-P: Immunohistochemistry (Paraffin) **Applications Key** 

IF-IC: Immunofluorescence (Immunocytochemistry)

H: human M: mouse R: rat Hm: hamster Mk: monkey Vir: virus Mi: mink C: chicken Dm: D. melanogaster **Cross-Reactivity Key** 

X: Xenopus Z: zebrafish B: bovine Dg: dog Pg: pig Sc: S. cerevisiae Ce: C. elegans Hr: horse

GP: Guinea Pig Rab: rabbit All: all species expected

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## **Limited Uses**

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